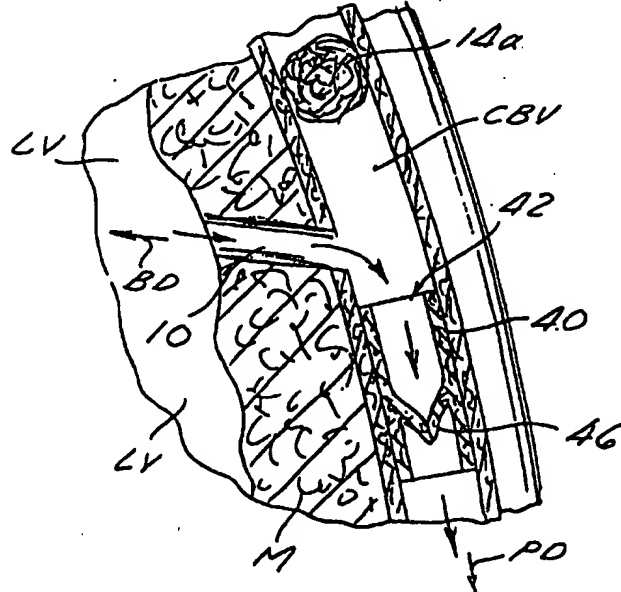




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<p>(54) Title: METHODS AND APPARATUS FOR TRANSMYOCARDIAL DIRECT CORONARY REVASCULARIZATION</p> <p>(57) Abstract</p> <p>This invention is methods and apparatus for direct coronary revascularization wherein a transmyocardial passageway (10) is formed between a chamber of the heart (LV) and a coronary blood vessel (CBV) to permit blood to flow therebetween. In some embodiments, the transmyocardial passageway (10) is formed between a chamber of the heart (LV) and a coronary vein (CV). The invention includes unstent transmyocardial passageways (10), as well as transmyocardial passageways (10) wherein protrusive stent devices (90) extend from the transmyocardial passageway (10) into an adjacent coronary vessel (CBV) or chamber of the heart (LV). The apparatus of the present invention includes protrusive stent devices (90) for stent of transmyocardial passageways (10), intraluminal valve devices (40) for valving of transmyocardial passageways (10), intra-cardiac valve devices (80) for valving of transmyocardial passageways (10), endogenous tissue valves (50) for valving of transmyocardial passageways (10), and ancillary apparatus for use in conjunction therewith.</p> 		

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5 METHODS AND APPARATUS FOR TRANSMYOCARDIAL
 DIRECT CORONARY REVASCULARIZATION

Field of the Invention

 The present invention pertains generally to medical
10 treatment methods and devices, and more particularly to
 methods and devices for transluminal direct coronary
 revascularization.

Background of the Invention

 Coronary artery disease continues to be one of the
15 leading causes of morbidity and mortality, throughout the
 world. The typical etiology of coronary artery disease is
 characterized by the build-up of atherosclerotic plaque
 within the coronary arteries. Such deposits of
 atherosclerotic plaque tend to fully or partially block the
20 flow of blood through the affected coronary arteries, and
 if untreated can result in myocardial ischemia, infarction
 and death.

 For many years, the traditional surgical treatment of
 coronary artery disease has been coronary artery bypass
25 surgery. In traditional coronary artery bypass surgery,
 the patient is generally anesthetized and placed on
 cardiopulmonary bypass. A thoracotomy is performed and the
 obstructed coronary blood vessels are exposed by surgical
 dissection. One or more segments of the patient's saphenous
30 vein or internal mammary artery is/are harvested for use as
 bypass graft(s). The harvested segment(s) of vein or
 artery is/are then anastomosed to the obstructed coronary
 artery(ies) to form bypass conduit(s) around the arterial
 obstruction(s). Such traditional coronary artery bypass
35 surgery is expensive, extremely invasive, and is associated

with significant operative and perioperative complications.

One alternative to traditional coronary artery bypass surgery is balloon angioplasty. In balloon angioplasty, a flexible guide catheter is percutaneously inserted into a peripheral artery (e.g., the femoral artery) and is transluminally advanced through the vasculature until the distal tip of the catheter is within an obstructed coronary artery. Thereafter, a balloon catheter is passed through the guide catheter and into the obstructive lesion. The balloon of the balloon catheter is inflated one or more times to dilate coronary artery in the region of the obstructive lesion. These balloon angioplasty procedures tend to be less expensive and less traumatic than traditional coronary artery bypass surgery. However, balloon angioplasty procedures of this type have been associated with a significant incidence of restenosis at the angioplasty site. The cause and mechanism of such restenosis continues to be the subject of ongoing study. However, such restenosis has generally been attributed to either a) an increase in the mass of the artery wall (e.g., neointima formation), b) a thickening of the artery wall without substantial change in its mass (e.g., vascular remodeling) and/or c) radial contraction of the balloon-dilated artery wall upon healing of cracks and fissures that have been created by the balloon dilation process.

Another alternative to traditional coronary artery bypass surgery is transluminal atheroectomy or ablation of the obstructive matter within the coronary artery. These transluminal atheroectomy or ablation procedures are performed by passing a catheter-mounted ablation apparatus through the vasculature to the site of the coronary obstruction. the catheter-mounted ablative apparatus is then utilized to cut, shave, sonicate, pulverize or otherwise ablate the obstructive matter from the lumen of the coronary artery. These atheroectomy or ablative

procedures must be performed with caution to avoid abrasion or damage to the artery wall, as such abrasion or damage can result in excessive scarring and subsequent reocclusion of the artery lumen. Furthermore, these atheroectomy or ablative procedures may, in some cases at least, be confounded by the need to meticulously contain and remove the severed fragments of obstructive matter in order to prevent such fragments of obstructive matter from escaping into the patient's circulatory system. Examples of such atheroectomy catheters and other catheter-mounted ablative apparatus are described in United States Patent Nos. 3,433,226 (Boyd), 3,823,717 (Pohlman, et al.), 4,808,153 (Parisi), 4,936,281 (Stasz), 3,565,062 (Kuris), 4,924,863 (Sterzer), 4B70,953 (Don Michael, et al.), 5,069,664 (Suess, et al.), 4,920,954 (Alliger, et al.) and 5,100,423 (Fearnot), as well as foreign patents/patent publications EP0347098A2 (Shiber), WO87-05739 (Cooper), WO89-06515 (Bernstein, et al.), WO90-0130 (Sonic Needle Corp.), EP316789 (Don Michael, et al.), DE 3,821,836 (Schubert), DE2438648 (Pohlman), and EP 0443256A1 (Baruch).

Other alternatives to traditional coronary artery bypass surgery have included minimally invasive endoscopic procedures which, ostensibly at least, can be performed through small (e.g., 1-3cm) incisions formed in the patient's chest wall, by insertion of a thoracoscope and associated operative instruments through such incisions. One such thoracoscopic coronary bypass procedure is described in United States Patent No. 5,452,733 (Sternan et al.). If perfected, these minimally invasive coronary artery bypass procedures may lessen the discomfort and length of recovery time experienced by patients who undergo such minimally invasive procedures vis a vis those who undergo traditional coronary artery bypass surgery. However, the performance of endoscopic surgical procedures of this type typically requires a great deal of operator

skill and training. Furthermore, as with traditional coronary artery bypass surgery, the patients on whom these thoroscopic procedures are performed are likely to undergo general anesthesia (with or without cardiopulmonary
5 bypass) and the creation of a pneumothorax due to the formation of full-thickness incision(s) in the chest wall. Thus, many of the drawbacks associated with traditional coronary artery bypass surgery, are also associated with these minimally invasive thoroscopic procedures.

10 Another previously described procedure for bypassing coronary artery obstructions utilizes a transmyocardial passageway (e.g., an interstitial tunnel formed in the muscular wall of the heart) to carry blood from the left ventricle of the heart to an obstructed coronary artery.
15 Such procedure, hereinafter generally referred to as "Transmyocardial Direct Coronary Revascularization" (TMDCR) is described in United States Patent Nos. 5,287,861 (Wilk), 5,409,019 (Wilk), and 5,429,114 (Wilk). The TMDCR methods described in these prior patents require that a catheter be
20 introduced into the obstructed coronary artery and advanced through the obstructive lesion. After the catheter has been advanced through the obstructive lesion, the distal tip of the catheter is directed toward the artery wall and an opening (i.e., a transmyocardial passageway) is formed
25 through the artery wall, through the adjacent myocardium, and into the chamber of the left ventricle. Also, in this previously described TMDCR method, a stent is required to be positioned within the transmyocardial passageway. Such intramyocardial stent is constructed to perform a one-way
30 valving function (i.e., to open and close the transmyocardial passageway in accordance with changes in the systolic-diastolic cardiac cycle). These TMDCR methods, previously described in United States Patent Nos. 5,287,861 (Wilk), 5,409,019 (Wilk) and 5,429,114 (Wilk),
35 may be difficult or impossible to perform in patients who

suffer from total or near total obstructions of a coronary artery, because of the necessary for advancing the catheter through the coronary artery obstruction to accomplish creation of the transmyocardial passageway at a location which is downstream of the coronary obstruction. Furthermore, because these previously described TMDCR methods require placement of a stent within the transmyocardial passageway, such procedures are necessarily associated with procedural complexities associated with measuring and precutting the stent to a precise length so that it fits within the transmyocardial passageway without protruding into the chamber of the left ventricle and/or the lumen of the coronary artery. Also, any stent which is positioned solely within the transmyocardial passageway may be subject to repetitive flexing and/or stressing as the myocardium undergoes its normal contraction and relaxation. Such repeated flexing and/or stressing of the intramyocardial stent may lead to unwanted migration, dislodgement or damage of the stent.

In view of the above-summarized shortcomings and complexities of the previously described TMDCR methods, there exists a need in the art for the development of improved TMDCR methods and associated apparatus which may be utilized without the need for cumbersome stenting of the transmyocardial passageway and/or implantation of one-way valving apparatus within the transmyocardial passageway. Also, there exists a need for the development of a new TMDCR methods which can be performed in patients who suffer from total or near total coronary artery occlusion(s), without the need for advancing a catheter through such coronary artery occlusion(s).

Summary of the Invention

The present invention provides new TMDCR methods, as well as certain valving devices which are usable in conjunction with these TMDCR methods.

i. TMDCR Procedures Using Coronary Vein

In accordance with the invention, there is provided a specific TMDCR method wherein a transmyocardial passageway is formed between a chamber of the heart (e.g., left ventricle) and a coronary vein. In this embodiment of the invention, blood may pass from the cardiac chamber, through the transmyocardial passageway, and into the coronary vein for the purpose of improving blood flow to the myocardium and/or to equalize or normalize pressures within the coronary venous vasculature by draining blood from the vein into the cardiac chamber. The coronary vein of this embodiment may be situated next to an obstructed coronary artery, and one or more secondary blood flow passageways may be created between the coronary vein and the adjacent artery, at site(s) which is/are downstream of the coronary artery obstruction. Also, the lumen(s) of the coronary vein and/or adjacent coronary artery may be blocked or embolized at appropriate positions to facilitate the flow of blood in the desired direction(s) through the man-made blood flow passageway(s), the coronary vein and/or the coronary artery. Additionally, one or more valving apparatus may be positioned within the coronary vein and/or within the cardiac chamber, to control or intermittently block the flow of blood through the transmyocardial passageway.

ii. TMDCR Procedures Using Unstented
Transmyocardial Passageway

In accordance with the present invention, there is provided a method for coronary re-vascularization wherein an unstented transmyocardial passageway (e.g., a puncture tract, bore, tunnel, or other passageway) is formed between a chamber of the heart (e.g., the left ventricle) and a coronary vessel (e.g., a) an endogenous coronary artery; b) an endogenous coronary vein; c) a man-made passageway which

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has been formed in the heart, and which leads to an endogenous coronary vein; d) a man-made passageway which has been formed in the heart, and which leads to an endogenous coronary artery; and/or e) a man-made passageway which has been formed in the heart between an endogenous coronary artery and an endogenous coronary vein). The unstented transmyocardial passageway(s) created in accordance with this embodiment of the invention may be utilized to improve perfusion of the myocardium by shunting blood from the chamber of the heart (e.g., left ventricle) into the coronary vessel (e.g., vein artery or man-made passageway), or may alternatively be utilized to equalize or normalize flow or pressure within the cardiac vasculature by draining blood from one or more cardiac vessels (e.g., vein, artery or man-made passageway), into the chamber of the heart.

iii. Valving Devices Positionable in Coronary Vessels

Still further in accordance with the present invention, there are provided several types of intraluminal valving apparatus which may be positioned within the lumen(s) of the coronary blood vessel(s) (i.e., artery, vein or man-made passageway) which intersect with the transmyocardial passageway, to intermittently block bloodflow, in at least one direction, through the transmyocardial passageway. These intraluminal valving devices generally comprise tubular bodies having at least one occluder member positioned therein, said occluder member(s) being alternately moveable between i) open position(s) whereby bloodflow is permitted to pass through the transmyocardial bloodflow passageway in a desired direction, and ii) closed position(s) whereby blood is prevented from flowing through the transmyocardial bloodflow passageway, in an undesired direction.

iv. Tissue Valves For TMDCR Passageway

Alternatively, the present invention also includes endogenous tissue valve(s) which are formed in the transmyocardial passageway to perform a desired one-way valving function whereby blood is permitted to flow through the transmyocardial bloodflow passageway in a first direction, but is prevented from backflowing or regurgitating in a second direction.

v. Intracardiac Valving Devices For TMDCR Passageway

Still further in accordance with the present invention, there are provided intracardiac valving devices which are mountable within a chamber of the heart (e.g., left ventricle) immediately adjacent to an opening into a transmyocardial passageway which extends from the cardiac chamber to a coronary vessel (e.g., artery, vein or man-made passageway). Such intracardiac valving device may be constructed such that it will open in response to hemodynamic pressure generated during systole and/or in response to mechanical contraction (i.e., shortening and thickening) of the myocardium during systole. When open, the intracardiac valving device permits blood to flow through the transmyocardial bloodflow passageway. Thereafter, the valving device may be constructed to close when diastolic pressures are present in the cardiac chamber or when the myocardium undergoes mechanical relaxation (i.e., lengthening and thinning during diastole. When closed, the valving device will prevent blood from backflowing or regurgitating from the transmyocardial bloodflow passageway, into the cardiac chamber.

iv. Protrusive Stents and Stented Grafts

For TMDCR Passageways

Still further in accordance with the present invention, there are provided stents and stented grafts which are positionable within the transmyocardial passageway, and which protrude into the adjacent coronary

vessel (e.g., vein, artery or man-made passageway). These protrusive stents and/or protrusive stented grafts may be self-expanding or pressure-expandable. Optionally, one or more valves or occluder members may be positioned within
5 such protrusive stents and/or stented grafts to facilitate valving or directed movement of bloodflow in accordance with the diastolic/systolic cardiac cycle.

Further objects and advantages of the present invention will become apparent to those skilled in the art
10 upon reading and understanding of the following detailed descriptions of preferred embodiments.

Brief Description of the Drawings

Figure 1 is a perspective view of a human heart showing the typical anatomical positioning of the coronary
15 arteries and coronary veins of the left heart.

Figure 1a is a partial cut-away sectional view of a human heart wherein a transmyocardial passageway has been created between the left ventricle and a coronary vein, in accordance with the present invention.

20 Figure 1b is a partial longitudinal sectional view through an obstructed coronary artery and adjacent coronary vein, showing a transmyocardial passageway of the present invention, extending between the chamber of the left ventricle and the coronary vein.

25 Figure 1c is a partial longitudinal sectional view through an obstructed coronary artery and adjacent coronary vein, showing a transmyocardial passageway of the present invention extending between the chamber of the left ventricle to the coronary vein, and a secondary bloodflow
30 passageway extending from the coronary vein to the adjacent coronary artery, downstream of the obstruction.

Figure 1d is a partial longitudinal sectional view through a portion of the myocardium of a human heart, adjacent the left ventricle, showing an alternative
35 embodiment of the present invention wherein a

transmyocardial bloodflow passageway extends from the chamber of the left ventricle to a secondary passageway which has been created between the obstructed coronary artery and the adjacent coronary vein. Figure 2 is a
5 longitudinal sectional view showing a first embodiment of an intravascular valving apparatus of the present invention operatively positioned within a coronary blood vessel (artery, vein or man-made passageway).

Figure 2a is a perspective view of the intravascular
10 valving apparatus of Figure 2.

Figure 2b is an elevational view of a variant of the intravascular valving apparatus shown in Figures 2 and 2a, wherein a bloodflow blocking bulkhead is formed on the upstream end of the apparatus.

15 Figure 3 is a longitudinal sectional view of a second embodiment of an intravascular valving apparatus of the present invention operatively positioned in a coronary blood vessel (artery, vein or man-made passageway).

Figure 3a is longitudinal sectional view showing
20 variant of the second intravascular valving apparatus embodiment shown in Figure 3, wherein two (2) separate valving apparatus are respectively positioned upstream and downstream of the junction between the transmyocardial bloodflow passageway and the coronary blood vessel (artery,
25 vein or man-made passageway).

Figure 3b is a longitudinal sectional view of another variant of the second intravascular valving apparatus embodiment shown in Figure 3, wherein three (3) valves are
30 valving of bloodflow through a transmyocardial bloodflow passageway and coronary blood vessel (artery, vein or man-made passageway).

Figure 4 is a longitudinal sectional view showing a third embodiment of an intravascular valving apparatus of
35 the present invention operatively positioned within a

coronary blood vessel (artery, vein or man-made passageway).

Figure 5 is a longitudinal sectional view showing an intracardiac valving apparatus of the present invention along with an optional retainer assembly (dotted lines) useable to mount such intracardiac valving apparatus on the inner wall of the heart.

Figure 5a is a perspective view of the intracardiac valving apparatus of Figure 4 having the optional retainer assembly affixed thereto.

Figures 6a and 6b are longitudinal sectional views of a human heart wherein a bloodflow passageway has been created between the left ventricle and a coronary blood vessel (artery, vein or man-made passageway), and a valving tissue valve has been created in the wall of the blood vessel, in accordance with the present invention.

Figures 7a-7b are longitudinal sectional views of a human heart wherein a blood vessel passageway has been created between the left ventricle and a coronary blood vessel (artery, vein or man-made passageway), and wherein an elastic suture has been positioned, in accordance with the present invention.

Figure 8a is a longitudinal sectional view showing a protrusive stent apparatus of the present invention implanted within a transmyocardial passageway and extending into a coronary blood vessel (e.g., artery, vein or man-made passageway).

Figure 8b is a longitudinal sectional view showing an alternative embodiment of the protrusive stent apparatus shown in Figure 5a.

Figure 8c is a longitudinal sectional view showing another alternative embodiment of the protrusive stent apparatus shown in Figure 5a, having an optional tubular covering and/or optional valve(s) incorporated therein.

Detailed Description of the Preferred Embodiments

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The following detailed description and the accompanying drawings are provided for purposes of describing and illustrating presently preferred embodiments of the invention only, and are not intended to limit the scope of the invention in any way.

Upon making reference to the accompanying figures, it will be noted that many of the figures include showings of human cardiovascular anatomy. The various anatomical structures shown in the figures are labeled in accordance with the following legend:

	AO	Aorta
	CBV	Coronary Blood Vessel
		(artery, vein or
		man-made passageway)
15	CA	Coronary Artery
	CAL	Coronary Artery Lumen
	CV	Coronary Vein
	CVL	Coronary Vein Lumen
	IVC	Inferior Vena Cava
20	SVC	Superior Vena Cava
	LV	Left Ventricle
	RV	Right Ventricle
	IVS	Intraventricular Septum
	M	Myocardium

25 i. TMDCR Method Utilizing Coronary Vein

With reference to Figures 1-4, the present invention includes methods for improving perfusion of regions of the myocardium M which are ischemic or otherwise affected by the existence of an obstruction OB within a coronary artery CA, by forming a transmyocardial passageway 10 which extends from a chamber of the heart, (e.g., left ventricle LV), to a coronary vein CV.

In some embodiments of this method, the transmyocardial passageway 10 will simply provide a flow of blood from the chamber of the heart and into the coronary

vein CV, such that the blood will pass in retrograde fashion through the coronary vein CV to perfuse the ischemic portion of the myocardium through the coronary vein, as shown in Figure 1b.

5 In other embodiments of the invention, a secondary bloodflow passageway 12 may be created between the coronary vein CV into which the transmyocardial passageway 10 extends and the obstructed coronary artery CA, at a location which is downstream of the obstruction OB, as
10 shown in Figure 1c. The formation of this secondary bloodflow passageway 12 allows blood from the chamber of the heart (e.g., the left ventricle LV) to initially flow through the transmyocardial passageway 10, through a segment of the coronary vein lumen CVL, through the
15 secondary bloodflow passageway 12, and into the coronary artery lumen CAL, at a location downstream of the coronary artery obstruction OB, as shown in Figure 2b. The secondary bloodflow passageway 12 which extends between the coronary vein CV and the coronary artery CA may optionally
20 be stented or internally supported by a stent, sleeve or coating (e.g., a polymer coating) to maintain patency of the secondary passageway 12.

In at least some applications, the coronary vein lumen CVL may be purposely blocked (e.g., ligated, embolized,
25 fused, welded, clamped, etc.) at site(s) upstream and/or downstream of the transmyocardial passageway 10. As shown in Figure 1b, when the transmyocardial passageway 10 formed for the purpose of shunting oxygenated blood into the coronary vein lumen CVL, a proximal embolization member 14a
30 may be positioned within the coronary vein lumen CVL, immediately upstream of transmyocardial passageway 10, to ensure that the shunted blood will flow, in the desired retrograde direction through the coronary vein CV. Similarly, as shown in Figure 1c, when a secondary
35 bloodflow passageway 12 is formed to carry the oxygenated

blood from the coronary vein lumen CVL into the coronary artery lumen CAL, downstream of the obstruction OB, a distal embolization member 14b may be positioned within the coronary vein lumen CVL immediately downstream of the secondary bloodflow passageway 12, to divert the flow of blood through the secondary bloodflow passageway 12.

Examples of methods for forming the optional secondary bloodflow passageway(s) 12 between the coronary vein CV and coronary artery CA are described in United States Provisional Specification Nos. 60/005,164, filed October 13, 1995 and 60/010,614 filed February 2, 1996, the entire disclosures of which are expressly incorporated herein by reference.

The proximal embolization member 14a and/or distal 14b embolization member may comprise any suitable type of lumen blocking matter or apparatus, examples of which are the embolization coils described in United States Patent Nos. 5,382,260 (Dormandy, Jr. et al.), 5,108,407 (Geremia et al.), and 5,256,146 (Ensminger, et al.). Alternatively, the coronary vein lumen CVL may be closed off at the sites of the proximal 14a and/or distal 14b embolization members by any suitable alternative means, such as clamping, clipping, ligating, fusing, welding or adhesively conjoining the inner walls of the coronary vein lumen CVL so as to provide the desired blocking of bloodflow therethrough.

Figure 1d shows an alternative embodiment of the method of the present invention wherein a secondary bloodflow passageway 12 of the above-described type has been created between the coronary vein CV and coronary artery CA, and wherein the transmyocardial bloodflow passageway 10a extends from the chamber of the heart (e.g., left ventricle) such secondary bloodflow passageway 12.

ii. TMDCR Methods Employing Non-Stented Transmyocardial Passageway

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The present invention also includes alternative TMDCR methods wherein a transmyocardial passageway 10 is formed between a chamber of the heart and a coronary vessel (i.e., a) an endogenous coronary vein, b) an endogenous coronary artery, c) a man-made passageway in the heart which connects to an endogenous coronary vein; d) a man-made passageway in the heart which connects to an endogenous coronary or e) a man-made passageway which extends between an endogenous coronary artery and an endogenous coronary vein), and such transmyocardial passageway 10 is allowed to remain non-stented (e.g., devoid of any stent or internal support member positioned therewith).

The utilization of a non-stented transmyocardial passageway 10 in accordance with this embodiment of the present invention eliminates the need for precise measurement, precutting to length and insertion of a stent apparatus within the transmyocardial passageway 10, as is required of the previous TMDCR method described in United States Patent Nos. 5,287,861 (Wilk), 5,409,019 (Wilk) and 5,429,114 (Wilk). When the non-stented transmyocardial passageway 10 of the present invention is intended to provide bloodflow from the chamber of the heart (e.g., left ventricle) into the coronary vessel (e.g., vein, artery or man-made passageway), the non-stented transmyocardial passageway 10 must remain open during systolic contraction of the myocardium. If the non-stented passageway 10 is permitted to substantially occlude or close-off during systolic contraction of the myocardium, such could prevent or deter the desired blood flow from passing through the transmyocardial passageway 10. In this regard, in embodiments of the invention which utilize the non-stented transmyocardial passageway 10, it may be desirable to debulk, core or otherwise enlarge the diameter of the passageway 10 during it's formation so as ensure that the passageway 10 will remain patent and open, even during

systolic contraction of the myocardium. Such coring, debulking or other enlargement of the passageway 10 may be accomplished by any suitable means, including the use of a hollow coring needle, laser, electrosurgical probe, or
5 other tissue removing/ablating device capable of debulking and removing tissue so as to create a transmyocardial passageway 10 of the desired diameter.

Also, it will be appreciated that the non-stented transmyocardial passageway preferably should not fill-in
10 with granulation tissue or otherwise close-off as a result of any scarring or healing process of the myocardium. In this regard, the coring, de-bulking or other enlargement of the non-stented passageway 10 and/or the continuing passage of blood, therethrough, may be sufficient to prevent or
15 deter such scarring or natural closing of the non-stented passageway 10. However, in applications wherein scarring or natural closing of the non-stented passageway 10 is a potential problem, it may be desirable to cauterize, heat, chemically treat or coat the walls of the non-stented
20 passageway to prevent or deter blocking of such passageway by scarring or ingrowth of the myocardial tissue.

iii. Valving Apparatus Positionable in the Coronary Vessel(s) to Prevent Backflow Into the Transmyocardial Bloodflow Passageway

25 In many embodiments of the invention, the transmyocardial passageway 10, 10a may function in it's intended manner without the inclusion of any valving apparatus, for intermittently blocking the flow of blood therethrough. However, in at least some applications, it
30 may be desired to prevent the backflow of blood through the transmyocardial passageway 10, 10a during certain phase(s) of the cardiac cycle when the relative hemodynamic pressures would tend to cause such backflow. In this regard, the present invention includes intravascular
35 valving apparatus 20, 30, 31, 33, 40, examples of which are

shown in Figures 2-4. These intravascular valving apparatus 20, 20, 31, 33, 40 are positionable within the lumen of the coronary blood vessel CBV (e.g., vein, artery or man-made passageway), and operate to prevent backflow of blood into
5 the transmyocardial bloodflow passageway 10, 10a.

In general, each of the intravascular valving apparatus 20, 30, 31, 33, 40 of the present invention comprise a radially expandable cylindrical or tubular body which is transluminally advanceable into the lumen of the
10 coronary blood vessel CBV (e.g., artery, vein or man-made passageway), and which is then radially expandable so as to become implanted at a location which is adjacent or near to the intersection of that coronary vessel CBV with a transmyocardial bloodflow passageway 10, 10a. The valving
15 apparatus 20, 30, 31, 33, 40 has an axial bore 24, 34, 42 through which blood may pass as it flows through the lumen of the coronary blood vessel CBV or secondary passageway 12 in which the apparatus 20, 30, 31, 33, 40 is positioned. One or more occluder members 26, 36, 46 are formed within
20 the apparatus 20, 30, 31, 33, 40. Such occluder member(s) 26, 36, 46 are alternately moveable between a first (e.g., open) position whereby blood is permitted to flow from the transmyocardial bloodflow passageway into the coronary blood vessel CBV or secondary passageway 12, and a second
25 (e.g., closed) position whereby blood is prevented or deterred from backflowing or regurgitating from the coronary blood vessel CBV or secondary passageway 12, into the transmyocardial bloodflow passageway.

Individual embodiments of the intravascular valving
30 apparatus 20, 30, 31, 33, 40 are described in more detail herebelow. It will be appreciated, however, that each of the intravascular valving apparatus 20, 30, 31, 33, 40 of the present invention offer advantages over the intramyocardial stenting/valving apparatus described in
35 United States Patent Nos. 5,248,861 (Wilk), 5,409,019

(Wilk) and 5,429,144 (Wilk) in that they are operatively situated entirely within the lumen of the coronary blood vessel CBV of secondary passageway 12 and do not extend into the transmural passage way (e.g., the first passageway 10, 10a) which emanates from the chamber (e.g., left ventricle) of the heart. In this regard, the valving apparatus 20, 30, 31, 33, 40 of the present invention do not require precise measurement or precise cutting-to-length, as is purportedly required of the intramyocardial stenting/valving apparatus described in U.S. Patent Nos. 5,248,861 (Wilk), 5,409,019 (Wilk) and 5,429,144 (Wilk).

It is desirable that the valving apparatus 20, 30, 31, 33, 40 of the present invention be initially disposable in a first radially compact diameter which is small enough to be mounted upon or inserted into an intravascular delivery catheter. Such intravascular delivery catheter, having the valving apparatus 20, 30, 31, 33, 40 mounted thereon or therewithin, is transluminally passable through the vasculature and into the lumen of the coronary blood vessel CBV wherein the apparatus 20, 30, 31, 33, 40 is to be implanted. Thereafter, the apparatus 20, 30, 31, 33, 40 is radially expanded (by self-expansion or pressure-expansion) to a second radially expanded diameter, wherein the outer surface of the apparatus 20, 30, 31, 33, 40 frictionally engages the surrounding wall of the coronary blood vessel CBV such that the apparatus 20, 30, 31, 33, 40 is thereby implanted and retained in a stationary position. When the valving apparatus 20, 30, 31, 33, 40 is so implanted within the coronary blood vessel CBV, blood may flow through the axial bore 24, 34, 42 of the apparatus 20, 30, 31, 33, 40, as described in more detail herebelow.

It is to be appreciated that the valving apparatus 20, 30, 31, 33, 40 may be either self-expanding or pressure-expandable. In this regard, if the valving apparatus 20, 30, 31, 33, 40 is "self-expanding", the cylindrical body of

the apparatus 20, 30, 31, 33, 40 may be formed of a shape memory alloy or resilient material (e.g., spring metal) which is inherently biased to it's second radially expanded diameter. Alternately, in embodiments wherein the valving apparatus 20, 30, 31, 33, 40 is "pressure-expandable", the cylindrical body of the apparatus 20, 30, 31, 33, 40 may be formed of plastically deformable material which is initially formed it's first radially compact diameter, and which may be pressure deformed to it's second radially expanded diameter by the exertion of outward force from an internally positioned balloon or other radial expansion device.

It is to be further appreciated that the potential useability and applicability of the intravascular valving apparatus 20, 30, 31, 33, 40, 50 described herebelow is not limited only to uses in connection with the improved TMDCR methods of the present invention, but may also be useable as a modification of the previously described TMDCR methods, such as those of United States Patent Nos. 5,287,816 (Wilk), 5,409,019 (Wilk), and 5,429,144 (Wilk).

a. Intravascular Valving Apparatus-First Embodiment:

Figures 2, 2a and 2b show a first embodiment of an intravascular valving apparatus 20 which is positioned within the lumen of a coronary blood vessel CBV (artery, vein or man-made passageway), at a location which is adjacent it's intersection with the transmyocardial passageway 10. This embodiment of the valving apparatus 20 has a cylindrical body having an axial bore 24 which extends longitudinally therethrough, and a side aperture 22 formed in the sidewall thereof. The side aperture 22 is preferably the same size or larger than the diameter of the adjacent end of the transmyocardial passageway 10, such that blood flowing from the cardiac chamber (e.g., left ventricle LV) through the transmyocardial passageway 10 will pass directly through the side aperture 22 and into

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the bore 24 of the valving apparatus 20. An occluder member 26, such as a hinged obturator or pliable elastomeric leaflet is affixed to the cylindrical body of the valving apparatus 20, and extends over and substantially blocks the side aperture 22 so as to prevent the flow of blood out of the side aperture 22. The occluder member 26 is alternately moveable between a first position wherein it blocks blood from flowing out of the side aperture 22, and a second position wherein it permits blood to flow into the bore 24 through the side aperture 22.

This first embodiment of the valving apparatus 20 may be implanted in the lumen of the coronary blood vessel CBV such that the side aperture 22 is in alignment with the adjacent end of the bloodflow passageway 10. During systolic contraction of the heart the relatively high pressure within the left ventricle will force the occluder member 26 to its second (open) position, allowing blood to flow from the left ventricle, through the transmural passageway 10, through the side aperture 22, through the bore 24 and into the lumen of the coronary blood vessel CBV in the perfusive direction PD, as shown. Thereafter, during systolic relaxation of the heart, the relatively low filling pressure within the left ventricle LV will draw the occluder member 26 to its first (closed) position whereby the occluder member 26 will prevent blood from regurgitating or moving in the backflow direction BD from the lumen of the coronary blood vessel CBV, out of the side aperture 22, and into the bloodflow passageway 10. In this manner the first embodiment of the valving apparatus 20 serves to facilitate efficient pumping of oxygenated blood from the left ventricle and into the lumen of the coronary blood vessel CBV, to improve the flow of oxygenated blood to an ischemic or blood-flow-deprived region of the myocardium M.

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As shown in Figure 2a, a closure member 21, in the nature of an end cap, may be formed on the upstream end of the apparatus 20 so as to completely or substantially block the flow of blood through the coronary blood vessel CBV and into the upstream end of the bore 24 of the apparatus 20. The optional inclusion of the end closure member 21 in the apparatus 20 may serve to obviate any need for the placement of a proximal embolization member 14a within the lumen of the coronary blood vessel CBV, upstream of the valving apparatus 20.

b. Intravascular Valving Apparatus-Second Embodiment

Figure 3 shows a second embodiment of the intravascular valving apparatus 30 which comprises a generally cylindrical body having an axial bore 34 extending longitudinally therethrough and a pair of occluder members 46 positioned therewithin, and a side aperture 32 formed in the cylindrical sidewall of the apparatus 30, behind the occluder members 36. Each occluder member 36 is affixed at least one point to the cylindrical body of the apparatus 30, and may comprise any suitable structure or openable and closeable passage, such as a self-sealing slit or hole, or a hinged leaflet or pliable elastomeric member. The occluder members 46 are alternately moveable between first positions wherein the occluder members 36 directly contact one another so as to prevent blood from backflowing in the backflow direction BD through the axial bore 34 of the apparatus 30, and second positions wherein the occluder members 36 move out of contact with one another such that blood may flow through the axial bore 34 of the apparatus 30 in the perfusion direction PD. The side aperture 32 is preferably as large as or larger than the diameter of the bloodflow passageway 10 which extends through the myocardium M from the left ventricle LV to the lumen of the coronary blood vessel CBV. This embodiment of the apparatus 30 is implanted in the

lumen of the coronary blood vessel CBV such that its side aperture 32 is directly aligned with the bloodflow passageway 10 so that blood may flow through the bloodflow passageway 10, into the axial bore 34 of the apparatus 30.

5 During systolic contraction of the heart the relatively high pressures created in the left ventricle LV will force blood to flow through the passageway 10 into the axial bore 34 of the valving apparatus 30. Such systolic bloodflow will move the occluder members 36 to
10 their second (i.e., open) positions, thereby allowing the blood to flow through the lumen of the coronary blood vessel in the perfusion direction PD. Thereafter, when the heart undergoes diastolic relaxation, the relatively low filling pressures created within the left ventricle LV will
15 draw the occluder members 36 to their first (ie. closed) positions, thereby preventing blood from regurgitating or backflowing out of the side aperture 32, in the backflow direction BD. In this manner, this second embodiment of the intravascular valving apparatus 30 serves to facilitate
20 efficient pumping of oxygenated blood from the left ventricle LV and through the lumen of the coronary blood vessel CBV, in order to provide improved bloodflow to an ischemic or blood-flow-deprived region of the myocardium M.

Optionally, secondary occluder members 38 may be
25 formed or mounted within the bore 34 of the apparatus 30, upstream of the side opening 32. These optional secondary occluder members 38 may be of the same type and construction as the above-described downstream occluder members 36. If present, such additional occluder members
30 38 will assume their first (e.g., closed) position when the pressure of blood within the bore 34 of the apparatus 30 downstream of such secondary occluder members 38 is greater than the pressure of blood within the coronary blood vessel CBV upstream of the such secondary occluder member 38. In
35 this regard, the provision of such secondary occluder

members 38 within the apparatus 30 will obviate the need for placement of a proximal occlusion apparatus 14a within the lumen of the coronary blood vessel CBV upstream of the transmyocardial bloodflow passageway 10. The inclusion of
5 such secondary occluder members 38, or the alternative use of a proximal occlusion member 14a, will be of particular importance when the coronary blood vessel CBV is a coronary vein CV, due to the substantial difference between endogenous coronary venous blood pressures and those
10 pressures which will be created by systolic arterial bloodflow through the coronary vein, downstream of the transmyocardial bloodflow passageway 10.

Figure 3a shows one variant of the second embodiment wherein two (2) separate intravascular valving apparatus
15 31a, 31b are respectively positioned upstream and downstream of the transmyocardial bloodflow passageway. The above-described occluder members 36 are formed in the apparatus 31b which is positioned downstream of the transmyocardial bloodflow passageway 10 and the above-
20 described secondary occluder members 38 are formed within the apparatus 31a which is positioned upstream of the transmyocardial bloodflow passageway 10. In this manner, these separate intravascular valving apparatus 31a, 31b, will function in the same manner as the apparatus 30 shown
25 in Figure 3, when it is equipped with the optional secondary occluder members 38. However, it will be appreciated that these separate intravascular valving apparatus 31a, 31b do not have any side aperture 32, as does the device shown in Figure 3, and accordingly, will
30 obviate any need for correctly sizing an aligning such side aperture 32 with the transmyocardial bloodflow passageway 10.

Figure 3b shows another variant of the second embodiment wherein a single intravascular valving apparatus
35 33, in the nature of a tubular stent or tubular body, is

provided with three (3) separate valves 26, 36, 38 at locations which are a) at the junction of the transmyocardial passageway 10 and the coronary blood vessel CBV, b) upstream of the transmyocardial passageway 10 and c) downstream of the transmyocardial passageway 10, respectively. These valves 26, 36, 38 may comprise self-sealing pliable slit openings, elastomeric leaflets, hinged occluder members or any other suitable type of structure or apparatus which will intermittently open and close, to permit bloodflow in the desired direction therethrough. For example, in applications wherein it is desired for the transmyocardial passageway 10 to provide a flow of blood from the cardiac chamber into the coronary blood vessel CBV, the first valve 26 will operate to open during systole to permit blood to flow from the transmyocardial passageway 10 into the coronary blood vessel CBV, but will close during diastole to prevent backflow or regurgitation into the cardiac chamber. Similarly, the second (upstream valve 38 will close during systole to prevent backflow of blood through the proximal end opening of the valving apparatus 33. The third (downstream) valve 36 will open during systole to permit the desired flow of blood entering through the transmyocardial passageway 10, to continue on downstream through the coronary blood vessel CBV in the desired perfusion direction.

c. Intravascular Valving Apparatus-Third Embodiment

Figure 4 shows a third embodiment of the intravascular valving apparatus 40 which comprises a generally cylindrical body having an axial bore 42 extending longitudinally therethrough and a plurality of occluder members 46 formed therewithin. The cylindrical body and occluder members 46 of this third embodiment of the apparatus 40 are the same as those of the above described second embodiment, except that the cylindrical body of this third embodiment is devoid of any side aperture(s) or

openings in the cylindrical sidewall. In contrast to the above described second embodiment 30, this third embodiment of the apparatus 40 is implanted in the lumen of the coronary blood vessel CBV at a location which is downstream
5 of the junction between the coronary blood vessel CBV and the first bloodflow passageway 10.

It will be appreciated that the individual features and attributes of each of the above-described embodiments of valving apparatus 20, 30, 31, 33, 40 may be incorporated
10 into any or all of the other above-described valving apparatus 20, 30, 31, 33, 40 as feasible, to accomplish the desired hemodynamic bloodflow within the coronary vasculature.

iv. Intracardiac Valving Apparatus For Controlling
15 Bloodflow Through the Transmyocardial Passageway

Figures 5 and 5a show examples of intracardiac valving apparatus 80 which may be utilized to prevent backflow of blood through the transmyocardial passageway 10, or to otherwise control the flow of blood through the
20 transmyocardial passageway 10 in accordance with the systolic/diastolic cardiac cycle.

As shown, the intracardiac valving apparatus 80 is positionable within the cardiac chamber (e.g., left ventricle) immediately adjacent the opening of the
25 transmyocardial passageway 10 therein. The intracardiac valving apparatus 80 may comprise any suitable type of hinged, pliable or moveable occlusion member or self-sealing slit which will operate to intermittently block or unblock the flow of blood in at least one direction through
30 the transmyocardial passageway 10. In the embodiment shown in Figures 5, 5a, the intracardiac valving apparatus 80 comprises a generally annular body having a central aperture formed therein and an occluder member 81, such as a pliable elastomeric flap, mounted within the aperture.
35 The occluder member 81 will move, in relation to

hemodynamic bloodflow and/or contraction of the myocardium M, between an open position whereby blood is permitted to pass in at least one direction through the transmyocardial passageway 10, and a closed position whereby blood is prevented from flowing in at least one direction through the transmyocardial passageway 10.

The intracardiac valving apparatus 80 may be implanted within the cardiac chamber by any suitable surgical or non-surgical technique. Preferably, the intracardiac valving apparatus 80 is initially positioned within or upon a delivery catheter, and the delivery catheter is advanced through the coronary blood vessel CBV, and through the transmyocardial passageway 10. Thereafter, the intracardiac valving apparatus 80 is released or ejected from the delivery catheter, and is caused to radially expand to it's operative configuration. the expanded valving apparatus 80 is then retracted into abutting contact with the myocardial wall, as shown.

The intracardiac valving apparatus 80 may be attached to the myocardial wall by any suitable attachment such as hooks, sutures, adhesives or a retaining assembly which is operative to hold the intracardiac valving apparatus 80 in its desired fixed position upon the myocardial wall. One such retaining apparatus, shown in Figures 5 and 5a, comprises an annular retaining ring 82 which is positionable within the coronary blood vessel CBV and a plurality of elastomeric tether members 84 which extend between the retainer ring 82 and the intracardiac valving apparatus 80. In this manner, the elastomeric tethers 84 will resiliently draw the retaining ring 82 and intracardiac valving apparatus 80 toward one another, so as to hold the intracardiac valving apparatus 80 in fixed abutment with the myocardium M as shown.

In some embodiments of the intracardiac valving apparatus 80, the occluder member 81 will be designed to

move in response to changes in hemodynamic pressure, such that when the hemodynamic pressure within the cardiac chamber (e.g., left ventricle) exceeds that within the transmyocardial passageway 10, the occluder member 81 will
5 move to it's open position, and when the pressure within the transmyocardial passageway 10 exceeds that within the cardiac chamber (e.g., left ventricle) the occluder member 81 will move to it's closed position.

Alternatively, in other embodiments of the
10 intracardiac valving apparatus 80, the occluder member 81 may be designed to move in relation to contractile changes in the myocardial muscle. In these embodiments, the occluder member 81 will be mechanically linked or coupled to the body of the intracardiac valving apparatus 80 such
15 that, when the myocardium undergoes contraction (e.g., shortening and thickening), the occluder member 81 will be propelled to it's open position, and when the myocardium undergoes relaxation (e.g., lengthening and narrowing) the occluder member 81 will move to it's closed position.

20 In this manner, the intracardiac valving apparatus 80 of the present invention serves to control the desired bloodflow through the transmyocardial passageway 10, without the need for customizing or precise cutting-to-size of any intramyocardial stent, as has been described in the
25 prior art.

v. Tissue Valve for Preventing Backflow into
the Transmyocardial Bloodflow Passageway

An alternative to the use of the above-described intravascular valving apparatus 20, 30, 31, 33, 40 and/or
30 the intracardiac valving apparatus 80, is an endogenous tissue valve which may be formed within the transmyocardial passageway 10 or at either end thereof. For example, Figures 6a-6b show an endogenous tissue valve 50 which is formed at the junction of the transmyocardial bloodflow

passageway 10 and a coronary blood vessel CBV (e.g., artery vein or man-made passageway).

With reference to Figure 6a-6b, the endogenous tissue valve 50 may comprise one or more segment(s) 54 of the wall of the coronary blood vessel CBV, along with one or more tapered segment(s) of underlying myocardial tissue 52.

This endogenous tissue valve 50 is formed such that the segment(s) of blood vessel wall 54 and underlying portion(s) of myocardial tissue 52 will receive sufficient blood supply so as not to become necrotic or infarcted. The thickness and mass of the tissue valve 50 is preferably defined so that, when the heart undergoes systolic contraction the elevated pressure created within the left ventricle LV and transmyocardial bloodflow passageway 10 will force the tissue valve 50 to an open position, as illustrated in Figure 5a, thereby creating an opening 56 through which blood may flow into the lumen of the coronary blood vessel CBV, in the perfusion direction PD. Thereafter, when the heart undergoes diastolic relaxation the relatively low filling pressures within the left ventricle LV and transmyocardial bloodflow passageway 10 will allow the tissue valve 50 to return to a second or closed position, as illustrated in Figure 5b. When in such second or closed position, the tissue valve 50 will substantially or completely close off the transmyocardial bloodflow passageway 10, so as to prevent blood from backflowing or regurgitating in the backflow direction BD, from the lumen of the coronary blood vessel CBV into the transmyocardial bloodflow passageway 10.

The tissue valve 50 may be created by any suitable means, including a procedure whereby the tissue penetrating, cutting or boring device used to create the transmyocardial bloodflow passageway is provided with a tapered distal end having a configuration analogous to that of the inner edge(s) 55 of the wall segment(s) 54 so as to

form the desired tissue valve(s) or segment(s) when form the endogenous tissue valve 50, or by another catheter-based device which is equipped to form such tissue valve(s) or segment(s).

5 It will be appreciated that the tissue valve 50 may be formed in various configuration. For example, although the tissue valve 50 shown in Figures 6a and 6b hereof consists of a single flap, various alternative configurations may be utilized wherein multiple tissue protrusions, multiple
10 tissue flaps, or annularly tapered or funnel shaped tissue flaps are formed to perform the desired valving function. Any and all such configurations of endogenous tissue are intended to be included within the scope of the term "tissue valve" 50 as used herein.

15 vi. Elastic Closure for Preventing Backflow Into the Transmyocardial Bloodflow Passageway

An alternative to the mechanical valving apparatus 20, 30, 31, 33, 40 or endogenous tissue valve 50 is the elastic closure member 60, shown in Figure 7a and 7b.

20 The elastic closure member 6d may comprise one or more sutures formed of stretchable or elastic material such as latex or other elastomeric polymer materials. Such elastic closure member(s) 60 are preferably passed through adjacent portions of myocardial tissue next to the opening 66
25 between the transmyocardial bloodflow passageway 10 and the lumen of the coronary blood vessel CBV (or secondary bloodflow passageway 12).

The elastic closure member(s) 60 is the elastically biased to a retracted state whereby the closure member(s)
30 60 will draw the adjacent portions of myocardium M together so as to close off the opening 66 between the transmyocardial bloodflow passageway 10 and the lumen of the coronary bloodflow CBV, as shown in Figure 7b. Upon systolic contraction of the heart the relatively high
35 pressures created within the left ventricle LV and

transmyocardial bloodflow passageway 10 will cause the elastic closure member(s) 60 to stretch or expand, thereby forming opening 66 through which blood may flow from the transmyocardial bloodflow passageway 10 into the lumen of the coronary blood vessel CBV (or secondary bloodflow passageway 12) in the perfusion direction PD, as shown in Figure 7a.

Thereafter, when the heart undergoes diastolic relaxation the relatively low filling pressures within the left ventricle LV and transmyocardial bloodflow passageway 10 will allow the elastic closure member 60 to retract, thereby closing off the opening 66 and preventing blood from backflowing or regurgitating from the lumen of the coronary blood vessel CBV (or secondary bloodflow passageway 12) into the transmyocardial bloodflow passageway 10, in the backflow direction BD, as shown in Figure 7b.

It will be appreciated that the elastic closure member 60 may be installed in any suitable method, such as by way of an appropriate suturing or stapling device which operates to attach the elastic closure member 60 at its desired location. Such installation of the elastic closure member 60 may be accomplished by open surgical technique or by way of catheter-based, transluminal methodology. For example, a catheter having a suturing or stapling device positioned therewithin may be advanced to a position adjacent the opening 66. Thereafter, negative pressure or other suitable drawings means may be utilized to draw adjacent segments of the myocardial tissue, from either side of the transmyocardial passageway 10, into the catheter. Thereafter, the desired elastic closure member 60 may be penetrated and threaded through the adjacent sides of the myocardial tissue so as to form the desired elastic closure member 60, as shown.

vii. Protrusive Stents and Stented Grafts for Stenting of

the Transmyocardial Passageway

In accordance with another aspect of the invention shown in Figures 8a-8c, protrusive stents or stented grafts may be positioned within the transmyocardial passageway 10, and may extend into one or more adjacent coronary vessels including a) an endogenous coronary vein, b) an endogenous coronary artery, c) a man-made passageway in the heart which connects to an endogenous coronary vein, d) a man-made passageway in the heart which connects to an endogenous coronary artery and/or e) a man-made passageway which extends between an endogenous coronary vein and an endogenous coronary artery. As described more fully herebelow, the protrusive stent apparatus 90, 90a, 90b of the present invention may incorporate one or more valving apparatus to intermittently block or direct bloodflow in accordance with various stages of the systolic/diastolic cardiac cycle. Furthermore, such protrusive stent apparatus may optionally be covered or juxtapositioned to a tubular graft or sheath so as to form a discrete tubular passageway.

Figure 8a shows a non-valved, non-covered protrusive stent apparatus 90 of the present invention positioned partially within a transmyocardial passageway 10, and extending into the coronary vessel CV (e.g., artery, vein or man-made passageway) to which such transmyocardial passageway 10 extends. As shown, the protrusive stent apparatus 90 is curved or bent at the junction of the transmyocardial passageway 10 and the coronary vessel CV, and preferably extends into the coronary vessel CV in the desired bloodflow direction.

The protrusive stent apparatus 90 may be formed of any suitable material, such as wire mesh or other metal or polymeric material, and may be self-expanding or pressure-expandable.

Figure 8b shows a variant of the protrusive stent

apparatus 90a positioned partially within a transmyocardial passageway 10, extending through a coronary vein CV, through a secondary passageway 12, and into a coronary artery CA. As shown the protrusive stent apparatus 90a is
5 curved or bent at the junction of the secondary passageway 12 and the coronary artery CA and preferably extends into the coronary artery CA in the desired bloodflow direction.

Figure 8c shows alternative variations of the protrusive stent apparatus 90b wherein an optional tubular
10 covering 92 is formed on the protrusive stent 90b. Such optional covering 92 may be any suitable tubular covering such as woven polyester or expanded, sintered polytetrafluoroethylene (PTFE). Additionally, or alternatively, one or more valves such as hinged occluder
15 members or pliable elastomeric leaflets may be located within the protrusive stent apparatus 90b with or without covering 92, at locations L_1 and/or L_2 and/or L_3 to facilitate control and valving of bloodflow through the transmyocardial passageway 10, coronary vein CV, secondary
20 passageway 12 and/or coronary artery CA. It will be appreciated that embodiments of the protrusive valving apparatus 90b which incorporates such valves at locations L_1 and/or L_2 and/or L_3 may be provided with appropriate openings or apertures in any covering 92 formed thereon to
25 facilitate the desired inflow or outflow of blood at specific locations thereon.

These protrusive stent apparatus 90, 90a, 90b with or without the optional covering 92 and/or without the optional valves at locations L_1 and/or L_2 and/or L_3 offer
30 advantages over previously known intramyocardial stents in that they do not require precise cutting to length or precise positioning within the myocardial passageway 10. Indeed, the protrusive stent apparatus 90, 90a, 90b of the present invention are intended to protrude into a coronary
35 blood vessel CBV (e.g., artery, vein and/or man-made

passageway) and the length of the portion of the stent apparatus 90, 90a, 90b which extends into such coronary blood vessel CBV is typically not critical. In this regard, there will exist no need for custom-fitting or
5 precise precutting of the stent apparatus 90, 90a, 90b prior to implantation within the patient.

In embodiments where the stent apparatus 90, 90a, 90b is covered by a partial or complete tubular covering, such covering may be formed of any suitable material including
10 but not necessarily limited to polyester, woven polyester, polytetrafluoroethylene, expanded polytetrafluoroethylene, polyurethane; silicone, polycarbonate, autologous tissue and, xenograft tissue.

The foregoing invention has been described hereabove with reference to certain presently preferred embodiments and examples only. No effort has been made to exhaustively describe all possible embodiments and examples in which the invention may be practiced. Indeed, various additions, deletions, modifications and alterations may be made to the
15 above-described embodiments and examples without departing from the intended spirit and scope of the invention. Accordingly, it is intended that all such additions, deletions and modifications and alterations be included
20 within the scope of the following claims.

WHAT IS CLAIMED IS:

1. A method for transmyocardial coronary revascularization, said method comprising the step of:

5 a) creating a transmyocardial bloodflow passageway between a chamber of the heart and a coronary vein.

2. The method of Claim 1 wherein said passageway is formed such that blood will flow from the chamber of the heart, through the transmyocardial bloodflow passageway, and through the lumen of the coronary vein, in a retrograde direction, so as to perfuse said region of the myocardium.

3. The method of Claim 1 wherein said coronary vein is situated next to a coronary artery, and wherein said method further comprises the step of:

15 b) forming a fistulous connection between said coronary vein and said adjacent coronary artery, at a location which is downstream of said transmyocardial bloodflow passageway, such that blood may flow from the chamber of the heart, through said transmyocardial bloodflow passageway, through said vein, through said fistulous connection, and into the adjacent coronary artery so as to provide enhanced bloodflow through said coronary artery.

4. The method of Claim 3 wherein said fistulous connection is a secondary bloodflow passageway which extends from said coronary vein to said coronary artery.

5. The method of Claim 1, further comprising the additional step of:

30 b) blocking the lumen of the coronary vein at a location which is upstream of said transmyocardial bloodflow passageway.

6. The method of Claim 3 wherein said method further comprises the steps of:

35 blocking the lumen of the coronary vein at a location downstream of said fistulous connection.

7. The method of Claim 1 further comprising the step of:

b) placing an intraluminal valving apparatus within the lumen of the coronary vein, said intraluminal valving apparatus comprising at least one occluder member which is alternately deployable in:

- i) an open position whereby bloodflow is permitted to pass through the transmyocardial bloodflow passageway in a first direction; and,
- 10 ii) a closed position whereby blood is prevented from backflowing through said transmyocardial bloodflow passageway, in a backflow direction.

8. The method of Claim 7 wherein said transmyocardial bloodflow passageway is intended to provide a flow of blood from the chamber of the heart into said coronary vein, and wherein said first direction is the direction extending from said chamber of the heart to said coronary vein, and wherein said second direction is the direction extending from said coronary vein to the chamber of the heart.

9. The method of Claim 7 wherein said transmyocardial bloodflow passageway is intended to drain blood from the coronary vein into the chamber of the heart, and wherein said first direction is the direction extending from the coronary vein to the chamber of the heart, and said second direction is the direction extending from the chamber of the heart to the coronary vein.

10. The method of Claim 7 wherein the intravascular valving apparatus provided in step B is positioned downstream of the transmyocardial bloodflow passageway, and wherein said method further comprises the step of:

c) blocking the lumen of the coronary vein upstream of the transmyocardial bloodflow passageway.

11. The method of Claim 7 wherein the intravascular

valving apparatus provided in step B is positioned downstream of the transmyocardial bloodflow passageway, and wherein said method further comprises the step of:

- 5 c) placing a second intravascular valving apparatus within the lumen of the coronary vein, upstream of said transmyocardial bloodflow passageway.

12. The method of Claim 1 further comprising the step of:

- 10 c) forming an endogenous tissue valve which is alternately moveable between:

- i) an open position whereby bloodflow is permitted to pass from said transmyocardial bloodflow passageway and through the lumen of said coronary vein, in a perfusion direction; and,
- 15

- ii) a closed position whereby said tissue valve will prevent blood from flowing from the coronary vein into said transmyocardial bloodflow passageway, in a backflow direction.

20 13. The method of Claim 12 wherein said tissue valve is formed at the junction of the transmyocardial bloodflow passageway and the coronary vein.

25 14. The method of Claim 13 wherein the tissue valve comprises at least one segment of the coronary vein in combination with at least one underlying segment of myocardial tissue.

30 15. The method of Claim 14 wherein at least one segment of coronary vein and the at least one segment of underlying tapered segment of myocardial tissue which form said tissue valve are sized and configured such that, when systolic blood pressure is created within said transmyocardial bloodflow passageway, said tissue valve will move to its open position, and thereafter when diastolic blood pressure is present in said transmyocardial bloodflow passageway, said tissue valve will move to its

35

closed position.

16. The method of Claim 1 further comprising the step of:

5 connecting an elastic closure member to cardiac tissue on either side of said transmyocardial bloodflow passageway, said elastic closure member being alternately transitionable between:

- 10 i) a stretched configuration whereby said transmyocardial bloodflow passageway is opened to permit blood to flow from said transmyocardial bloodflow passageway into said coronary vein; and
- 15 ii) a retracted configuration whereby said transmyocardial bloodflow passageway is substantially blocked so as to prevent blood from backflowing from said coronary vein into said transmyocardial bloodflow passageway.

17. The method of Claim 16 wherein said elastic closure member comprises a suture which is formed of elastic material, said suture being threaded through said

20 myocardial tissue on opposite sides of said transmyocardial bloodflow passageway.

18. The method of Claim 1 further comprising the step of:

25 b) placing an intracardiac valving apparatus within the chamber of the heart, adjacent one end of said transmyocardial bloodflow passageway, said intracardiac valving apparatus being alternately deployable in:

- 30 i) an open position whereby bloodflow is permitted to pass through the transmyocardial bloodflow passageway in a first direction; and,
- 35 ii) a closed position whereby blood is prevented from backflowing through the transmyocardial bloodflow passageway, in a second direction, said second direction being opposite

said first direction.

19. The method of Claim 18 wherein said transmyocardial bloodflow passageway is intended to provide a flow of blood from the chamber of the heart to the coronary vein, and wherein said first direction is the direction extending from the chamber of the heart to the coronary vein, and said second direction is the direction extending from the coronary vein to the chamber of the heart.

20. The method of Claim 18 wherein said transmyocardial bloodflow passageway is intended to drain blood from the coronary vein into the chamber of the heart, and wherein said first direction is the direction extending from the coronary vein to the chamber of the heart, and said second direction is the direction extending from the chamber of the heart to the coronary vein.

21. The method of Claim 18 wherein the intracardiac valving apparatus provided in step b is attached to the wall of the chamber of the heart, and is positioned over the opening formed in the chamber of the heart by said transmyocardial bloodflow passageway.

22. The method of Claim 21 wherein said intracardiac valving apparatus is sutured to the wall of the chamber of the heart.

23. The method of Claim 21 wherein said intracardiac valving apparatus is adhered to the wall of the chamber of the heart.

24. The method of Claim 1 further comprising the step of:

b) placing a protrusive stent within said transmyocardial passageway, such that said protrusive stent extends into said coronary vein.

25. The method of Claim 24 wherein said protrusive stent is uncovered.

26. The method of Claim 24 wherein said protrusive

stent is at least partially covered.

27. The method of Claim 24 wherein said protrusive stent incorporates at least one valve to intermittently block blood flow, in at least one direction, through said transmyocardial passageway.

28. The method of Claim 27 wherein said valve is operative to permit blood to flow from said chamber of the heart through said transmyocardial passageway, and into said coronary vein, but will prevent blood from backflowing from said coronary vein into said transmyocardial passageway.

29. A method for transmyocardial direct coronary revascularization, said method comprising the steps of:

a) forming a transmyocardial bloodflow passageway from a chamber of the heart to a coronary blood vessel;

b) permitting blood to flow from the chamber of the heart, through said transmyocardial bloodflow passageway; and

c) into the coronary blood vessel, while said transmyocardial bloodflow passageway remains devoid of any stent positioned therewithin.

30. The method of Claim 29 wherein said blood vessel is selected from the group consisting of:

i) an endogenous coronary artery;

ii) an endogenous coronary vein;

iii) a man-made passageway which has been formed in the heart and which connects to an endogenous coronary vein;

iv) a man-made passageway which has been formed in the heart and which connects to an endogenous coronary artery; and

v) a man-made passageway which extends between an endogenous coronary artery and an endogenous coronary vein.

31. The method of Claim 29 wherein said coronary blood vessel is an endogenous coronary vein which is situated next to a coronary artery, and wherein said method further comprises the step of:

- 5 d) forming a second bloodflow passageway between said coronary vein and the adjacent coronary artery, at a location which is downstream of said transmyocardial bloodflow passageway.

32. The method of Claim 31 wherein said second
10 bloodflow passageway is a fistulous tract which extends between said coronary vein and said coronary artery.

33. The method of Claim 29 wherein it is intended for blood to flow in a first flow direction through said coronary blood vessel and wherein said method further
15 comprises the additional step of:

- d) blocking the lumen of the coronary blood vessel at a location which is upstream of said transmyocardial bloodflow passageway.

34. The method of Claim 31 wherein said method
20 further comprises the step of:

- d) blocking the lumen of the coronary vein downstream of said fistulous connection.

35. The method of Claim 29 further comprising the step of:

- 25 d) placing an intraluminal valving apparatus within the lumen of the coronary blood vessel, said intraluminal valving apparatus comprising at least one occluder member which is alternately deployable in:

 i) an open position whereby bloodflow is
30 permitted to pass from said transmyocardial bloodflow passageway and through the lumen of the coronary vein in a perfusion direction; and,

 ii) a closed position whereby blood is
35 prevented from flowing from the coronary vein into said transmyocardial bloodflow passageway,

in a backflow direction.

36. The method of Claim 35 wherein the intravascular valving apparatus of step C is positioned downstream of the transmyocardial bloodflow passageway, and wherein said
5 method further comprises the step of:

e) blocking the lumen of the coronary vein upstream of the transmyocardial bloodflow passageway.

37. The method of Claim 29 further comprising the step of:

10 c) forming an endogenous tissue valve which is alternately moveable between:

i) an open position whereby bloodflow is permitted to pass from said transmyocardial bloodflow passageway and through the lumen of
15 said coronary vessel, in a first direction; and,

ii) a closed position whereby said tissue valve will prevent blood from flowing from the coronary vein into said transmyocardial bloodflow passageway, in a second direction opposite said
20 first direction.

38. The method of Claim 37 wherein said tissue valve is formed at the junction of the transmyocardial bloodflow passageway and the coronary blood vessel.

39. The method of Claim 38 wherein the tissue valve
25 comprises at least one segment of the coronary blood vessel in combination with at least one underlying segment of myocardial tissue.

40. The method of Claim 37 wherein at least one segment of coronary blood vessel and at least one
30 underlying tapered segment of myocardial tissue which form said valving tissue valve are sized and configured such that, when systolic blood pressure is created within said transmyocardial bloodflow passageway said tissue valve will move to its open position, and thereafter when diastolic
35 blood pressure is present in said transmyocardial bloodflow

passageway, said tissue valve will move to its closed position.

41. The method of Claim 29 further comprising the step of:

5 connecting an elastic closure member to the myocardial tissue on either side of said transmyocardial bloodflow passageway, said elastic closure member being alternately transitionable between:

10 i) a stretched configuration whereby an opening is formed to permit blood to flow from said transmyocardial bloodflow passageway into said coronary vein; and

15 ii) a retracted configuration whereby said opening is substantially closed, thereby preventing blood from backflowing from said coronary vein into said transmyocardial bloodflow passageway.

20 42. The method of Claim 41 wherein said elastic closure member is a suture which is formed of elastic material and passed through said myocardial tissue on opposite sides of said transmyocardial bloodflow passageway.

25 43. The method of Claim 29 further comprising the step of:

 b) placing a protrusive stent within said transmyocardial passageway, such that said protrusive stent extends into said coronary vessel.

30 44. The method of Claim 43 wherein said protrusive stent is uncovered.

 45. The method of Claim 43 wherein said protrusive stent is at least partially covered.

35 46. The method of Claim 43 wherein said protrusive stent incorporates at least one valve to intermittently block blood flow, in at least one direction, through said

transmyocardial passageway.

47. The method of Claim 46 wherein said valve is operative to permit blood to flow from said chamber of the heart through said transmyocardial passageway, and into
5 said coronary vessel, but will prevent blood from backflowing from said coronary vein into said transmyocardial passageway.

48. An intraluminal valving apparatus which is operative to prevent blood from a backflowing from a
10 coronary blood vessel into a transmyocardial bloodflow passageway which extends from a chamber of the heart to said coronary blood vessel, said apparatus comprising:

a generally cylindrical body having an axial bore which extends longitudinally therethrough; and,
15 at least one occluder member positioned within said axial bore, said at least one occluder member being alternately moveable between:

i) an open position whereby systolic blood is permitted to pass from said transmyocardial
20 bloodflow passageway, through the lumen of the coronary blood vessel; and,

ii) a closed position whereby blood is prevented from backflowing from the lumen of the coronary blood vessel into the transmyocardial
25 bloodflow passageway.

49. The valving apparatus of Claim 48 wherein said generally cylindrical body is initially of a radially compact diameter so as to be transluminally advancable through the vasculature into said blood vessel, and is
30 subsequently expandable to a second radially expanded diameter wherein said cylindrical body will contact and engage the surrounding wall of said blood vessel.

50. The valving apparatus of Claim 49 wherein said cylindrical body is self-expanding.

35 51. The valving apparatus of Claim 49 wherein said

cylindrical body is pressure-expandable.

52. The valving apparatus of Claim 48 further comprising:

5 a side aperture formed in the cylindrical body of said apparatus, said side aperture being alignable with said transmyocardial bloodflow passageway such that blood from said transmyocardial bloodflow passageway may flow through said side aperture and into the axial bore of the valving apparatus. '

10 53. The valving apparatus of Claim 52 wherein said at least one occluder member is configured to close off said side aperture when in it's closed position, and further such that a subsequent increase in blood pressure within the transmyocardial bloodflow passageway will move said
15 occluder member to said open position, thereby reopening said side aperture.

54. The valving apparatus of Claim 53 wherein said at least one occluder member is positioned within the axial bore of the apparatus, at a location downstream of said
20 side aperture, such that systolic bloodflow which passes from the transmyocardial bloodflow passageway into the axial bore of the apparatus will force said occluder member to its open position, thereby causing the bloodflow to continue in the downstream direction, and the subsequent
25 creation of diastolic blood pressure is within the transmyocardial bloodflow passageway will move said occluder member to its closed position thereby preventing blood from backflowing out of said side aperture and into said transmyocardial bloodflow passageway.

30 55. The valving apparatus of Claim 53 further comprising:

a blocking member which closes off the axial bore of the apparatus, upstream of said side aperture.

56. The valving apparatus of Claim 48 further
35 comprising:

a secondary occluder member which closes off the axial bore of the apparatus, upstream of said side aperture.

5 57. The valving apparatus of Claim 48 wherein the apparatus is intended to be positioned within said coronary blood vessel at a location downstream of said junction between said blood vessel and said transmyocardial bloodflow passageway, and wherein:

10 said at least one occluder member which is configured to permit blood to flow in a perfusion direction through said axial bore, when said at least one occluder member is in it's open position, and to prevent blood from backflowing through said axial bore in a backflow direction, when said occluder member is
15 in its closed position.

58. A system comprising two of the valving apparatus of Claim 48, one of said valving apparatus being positionable within said coronary blood vessel upstream of said transmyocardial bloodflow passageway, and the other of
20 said valving apparatus being positionable within said coronary blood vessel downstream of said transmyocardial bloodflow passageway.

59. An intracardiac valving apparatus which is operative to control bloodflow through a transmyocardial
25 bloodflow passageway extending from a chamber of the heart to a coronary blood vessel, said intracardiac valving apparatus comprising:

a valve body having an opening formed therethrough, said valve body being positionable in
30 contact with the wall of the heart such that the opening of said valve body is in alignment with said transmyocardial bloodflow passageway;

at least one occluder member positioned within the opening of said valve body, said occluder member being
35 alternately moveable between:

i) an open position whereby blood is permitted to pass through said transmyocardial bloodflow passageway in a first direction; and,

5 ii) a closed position whereby blood is prevented from flowing through said transmyocardial passageway in at least a second direction opposite said first direction.

60. The intracardiac valving apparatus of Claim 59 wherein said apparatus further comprises:

10 means for holding said intracardiac valving apparatus in substantial fixed position against said wall of the chamber of the heart.

61. The intracardiac valving apparatus of Claim 60 wherein said means for holding comprises hooks.

15 62. The intracardiac valving apparatus of Claim 61 wherein said means for holding comprises sutures.

63. The intracardiac valving apparatus of Claim 61 wherein said means for holding comprises an adhesive.

20 64. The intracardiac valving apparatus of Claim 61 wherein said means for holding comprises a retainer assembly which engages the heart, and which exerts force upon said intracardiac valving apparatus to hold said intracardiac valving apparatus to hold said intracardiac valving apparatus in substantially fixed position.

25 65. The intracardiac valving apparatus of Claim 64 wherein said retainer assembly comprises:

30 a generally annular retainer ring having an aperture formed therein, said generally annular retainer ring positionable within the coronary blood vessel such that the aperture formed therein is in alignment with the transmyocardial passageway; and,

35 at least one elastomeric tether member having a first end connected to said intracardiac valving apparatus and a second end connected to said retainer ring, said elastomeric tether member being of a length

and resiliency which is sufficient to exert sufficient inward pressure upon said valving apparatus and said retainer ring to hold said valving apparatus and said retainer ring in substantially fixed positions, with the aperture of the valving apparatus and the aperture of the retainer ring being in alignment with said transmyocardial passageway.

66. A protrusive stent apparatus for stenting a transmyocardial passageway which extends from a chamber of the heart to a coronary blood vessel, said apparatus comprising:

a tubular body which is alternately configurable in:

- i) a radially collapsed configuration of a first diameter;
- ii) a radially expanded configuration of a second diameter, said second diameter being at least as large as the diameter of the transmyocardial passageway;

said protrusive stenting apparatus having a length which is longer than the length of the transmyocardial passageway, such that said apparatus may be positioned within said transmyocardial passageway from said cardiac chamber to said coronary blood vessel, with a portion of said apparatus protruding into said coronary blood vessel.

67. The protrusive stent apparatus of Claim 66 wherein the tubular body of said stent apparatus is self-expanding.

68. The protrusive stent apparatus of Claim 66 wherein the tubular body of said stent apparatus is pressure expandable.

69. The protrusive stent apparatus of Claim 66 wherein the tubular body of said stent apparatus is formed of material selected from the group of materials consisting

of:

metal;
polymeric material.

70. The apparatus of Claim 66 wherein said apparatus
5 further comprises:

a tubular covering formed on said stent.

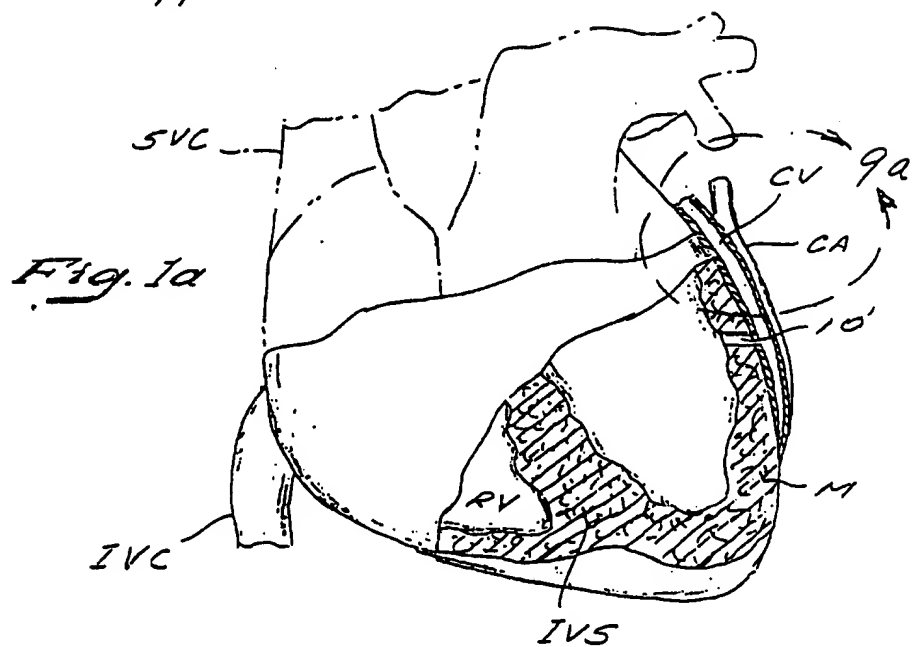
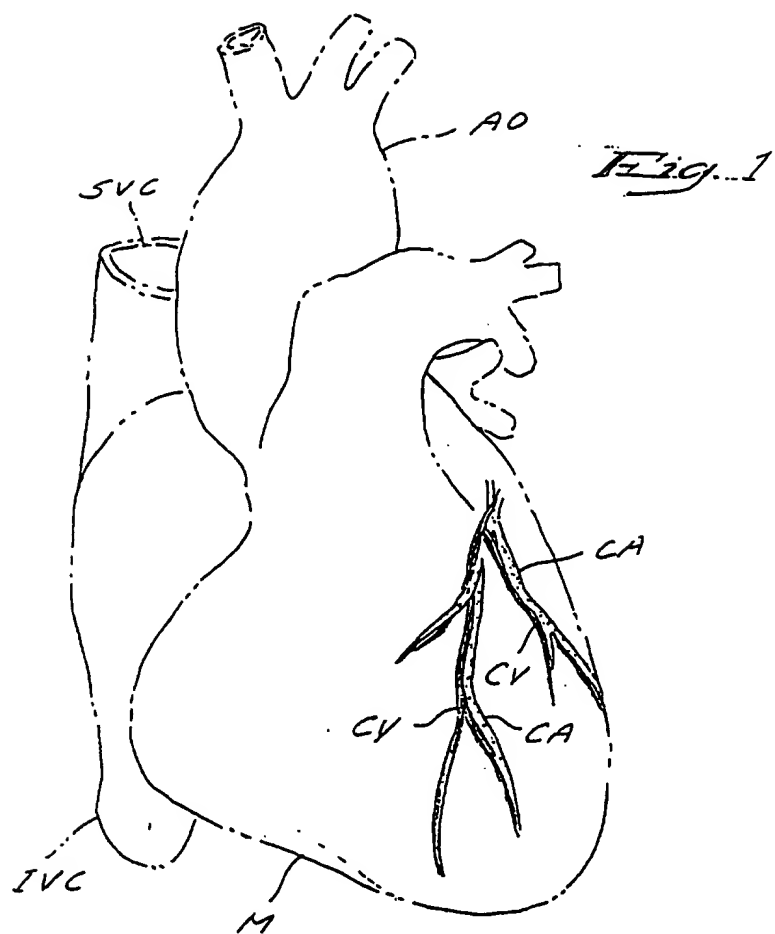
71. The apparatus of Claim 70 wherein said tubular
covering is formed of a material selected from the group of
materials consisting of:

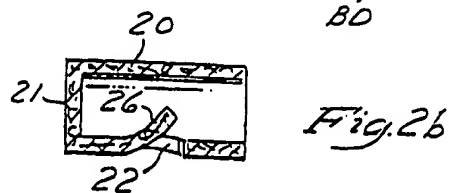
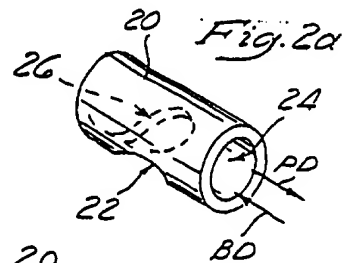
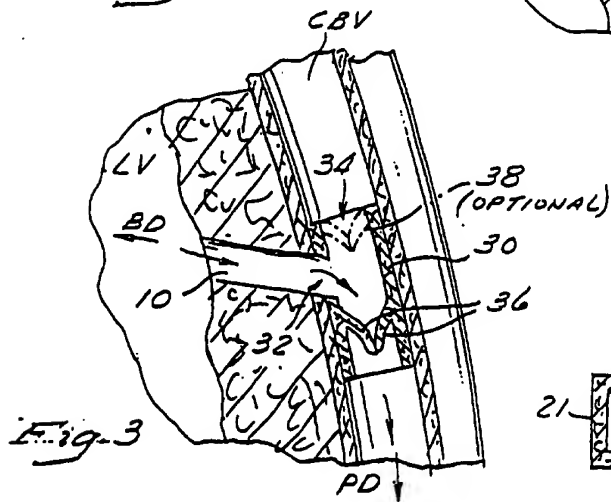
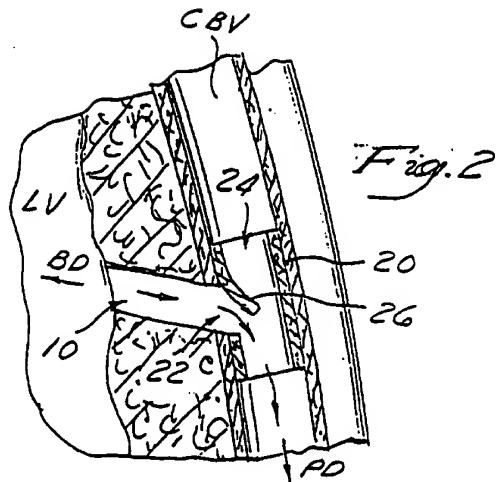
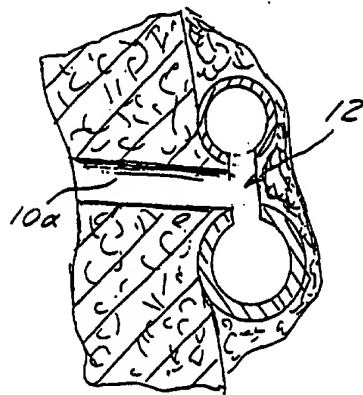
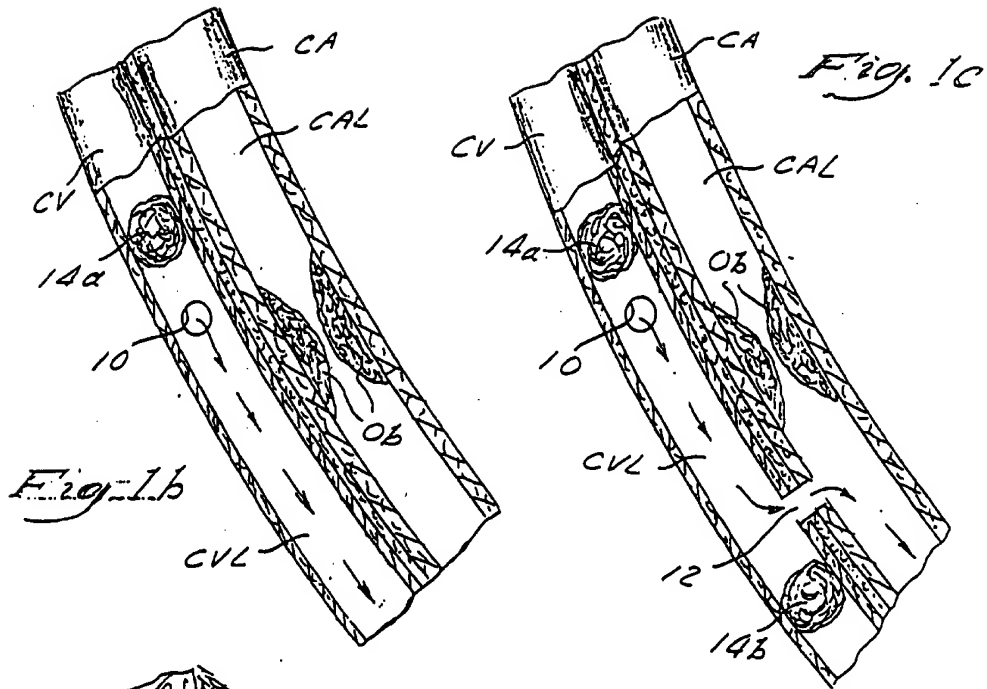
10 polyester;
woven polyester;
polytetrafluoroethylene;
expanded polytetrafluoroethylene;
polyurethane;
15 silicone;
polycarbonate;
autologous tissue; and,
xenograft tissue.

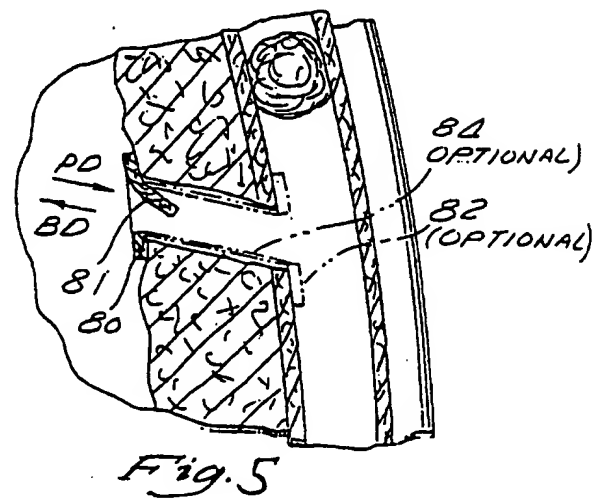
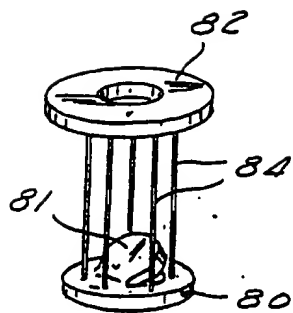
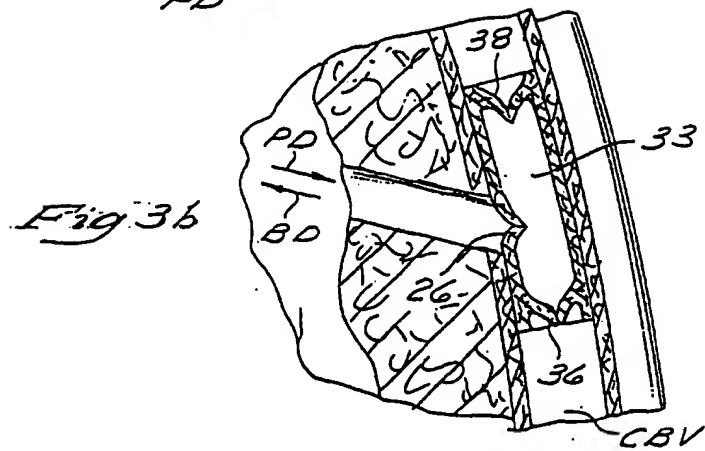
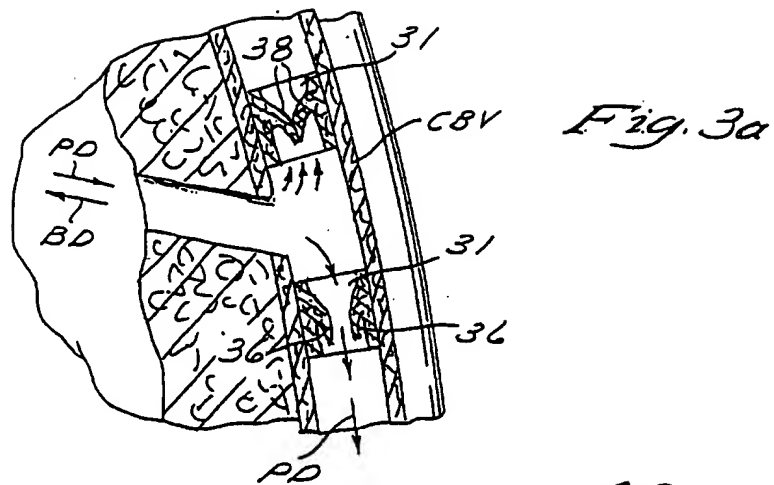
72. The apparatus of Claim 66 wherein said apparatus
20 further comprises:

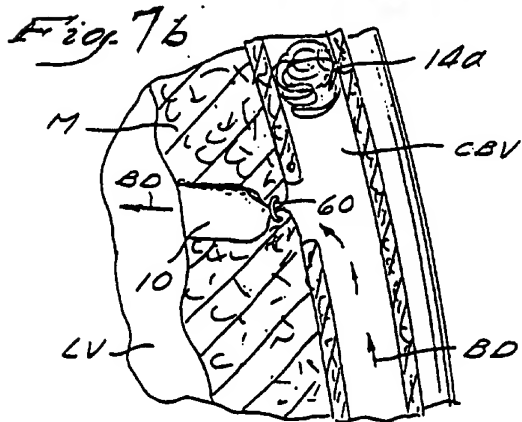
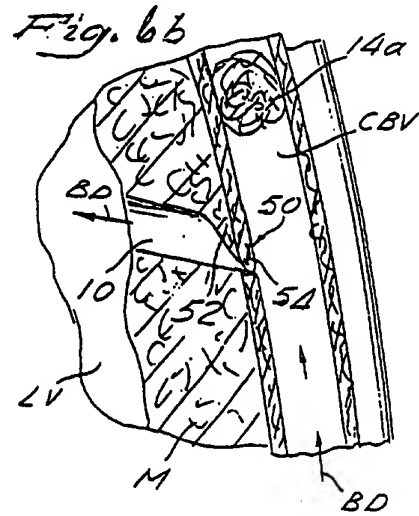
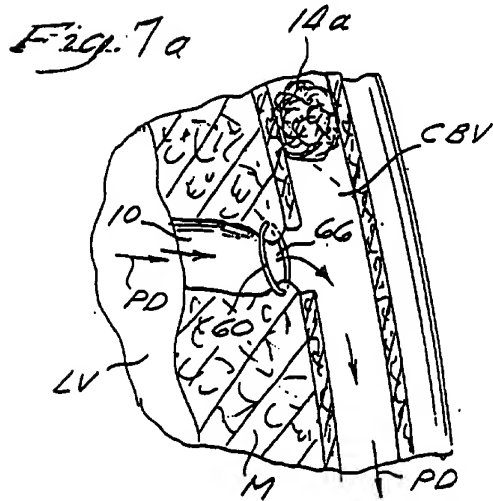
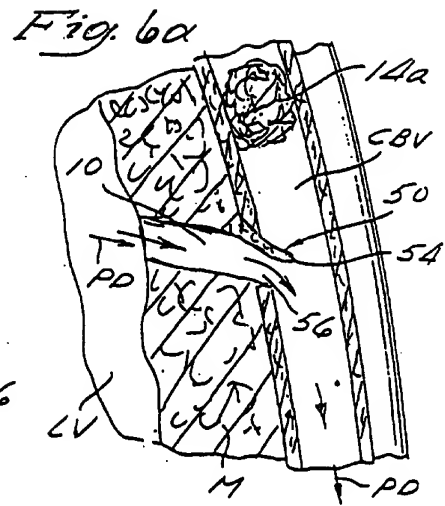
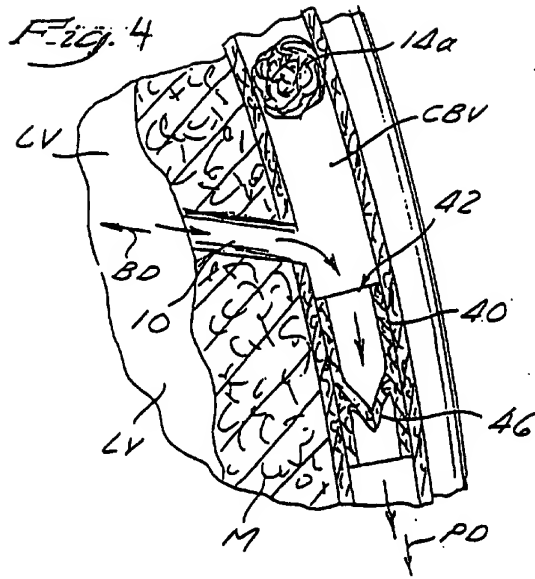
at least one valve positioned within said tubular
body to control bloodflow therethrough.

25









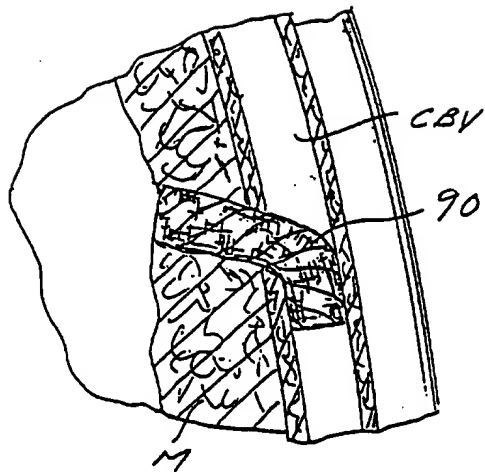


Fig. 8a

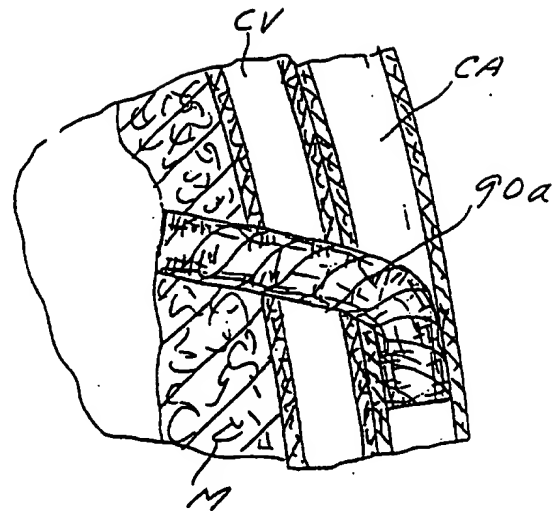


Fig. 8b

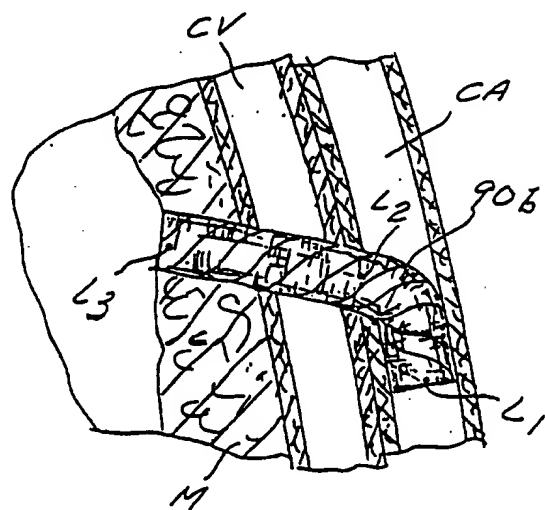


Fig. 8c

INTERNATIONAL SEARCH REPORT

International application No.

PCT/US97/14801

A. CLASSIFICATION OF SUBJECT MATTER

IPC(6) :A61B 19/00

US CL :128/898

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 128/898; 600/16; 606/7, 14; 623/1, 2

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS

Search Terms: transmyocordi? and passageway#

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X --- Y	US 5,429,144 A (WILK) 22 February 1994, entire document.	27, 30, 41, 43-51, 57, 59, 60, 64, 66-68, 70, 72 ----- 61, 62, 63, 69, 71
A	US 5,500,014 A (QUIJANO et al) 19 March 1996, entire document.	1-72
A	US 5,389,096 A (AITA et al) 14 February 1995, entire document.	1-72

☐ Further documents are listed in the continuation of Box C.☐ See patent family annex.

* Special categories of cited documents:	* T	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principles or theory underlying the invention
* A		document defining the general state of the art which is not considered to be of particular relevance
* B		earlier document published on or after the international filing date
* L		document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
* O		document referring to an oral disclosure, use, exhibition or other means
* P		document published prior to the international filing date but later than the priority date claimed
	* X	document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
	* Y	document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
	* A	document member of the same patent family

Date of the actual completion of the international search

08 DECEMBER 1997

Date of mailing of the international search report

23 DEC 1997

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